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**Michael Yeadon: Corona vaccines batch scandal | www.kla.tv/22390.mp4 Author: Dr. Michael Yeadon**

## The Standard of Pharmaceutical Consistency

**Pharmaceutical manufacturing is an industry defined by its ability to achieve consistent high quality, purity, and reproducibility across billions of doses. Global companies like Pfizer and Janssen are experts in this field, having manufactured vast quantities of tablets, capsules, and injections for decades. The production of these medical products is a complex, two-step process: first, the creation of the active drug substance, such as mRNA or DNA, and second, the formulation of that substance into a finished drug product. This entire journey is governed by strict Good Manufacturing Practice (GMP) regulations and overseen by medical regulators who must approve every specific production plan. Because of these incredible levels of control, every vaccine lot is expected to contain effectively the same material, regardless of the date or location of its manufacture. Under normal circumstances, a random draw of vaccine lots from a reporting system should show very similar performance and adverse event profiles from batch to batch.**

## The Discovery of the "Hot Lot" Phenomenon

**Independent analyses of the Vaccine Adverse Event Reporting System (VAERS) database have uncovered a startling deviation from these manufacturing expectations. While one would expect a uniform scattering of adverse events across all states and vaccine lots, the data shows that approximately 90% of adverse events are concentrated in less than 10% of the batches. This pattern is unprecedented when compared to historical data for other injectable products. For example, decades of data on influenza vaccines, covering over 22,000 lots, show a consistent and low level of serious adverse events, averaging only about four per lot. These rare events essentially represent "background noise" or the natural health fluctuations of a large population, rather than a product defect. However, the COVID-19 vaccine lots display a completely different and highly alarming profile.**

## Extreme Variability and Mathematical Impossibility

**The level of toxicity and variability observed in COVID-19 vaccine lots is described as being orders of magnitude higher than anything seen with the flu vaccine. While some COVID-19 lots show very few side effects, others are associated with hundreds or even thousands of serious adverse events. A specific example is lot EN6201, which is linked to approximately 600 serious adverse event reports, including life-threatening conditions and hospitalizations. Mathematically, it is impossible for such extreme differences to occur if the manufacturing process were truly producing identical material. Small manufacturing**

**fluctuations cannot explain why one lot results in two reports while another results in 5,000. This level of variation indicates that the material in the highly toxic lots is fundamentally different from the material used in clinical trials or in the safer batches. Furthermore, preliminary studies suggest that these differences are not driven by the size of the lot, as no relationship has been found between the number of doses in a batch and the number of adverse events it generates.**

**The Case for Intentionality**

**The precision of modern pharmaceutical companies makes it highly unlikely that these toxic variations are accidental. These companies are professional outfits that monitor real-time VAERS data as their "window onto the world," meaning they are fully aware of which batches are causing harm. The fact that they have not halted distribution despite this knowledge suggests that the variation is, at minimum, accepted. Additionally, the theory that these "hot lots" are the result of accidental degradation or "cold chain" issues is scientifically weak. Typically, when a biological product like mRNA degrades, it loses its functional activity rather than acquiring extreme, new toxicity. Most importantly, this pattern of high variability and extreme toxicity is observed across three different manufacturers and two separate technologies—both mRNA and DNA-based vaccines. It is virtually impossible for three different companies to experience the same "mishap" using different technologies simultaneously.**

**Suppression of Information and Psychological Warfare**

**The public has been prevented from understanding these risks due to a coordinated effort by Big Tech and major media outlets. Companies like Google, Facebook, and YouTube have censored and deplatformed qualified scientists who disagree with official public health narratives. This control over information ensures that the public only hears a one-sided, and arguably misleading, description of the situation. Qualified professionals like Dr. Michael Yeadon, Dr. Robert Malone, and Dr. Peter McCullough have been barred from mainstream media studios to prevent them from "destroying" the official story with their data and expertise. This environment of fear and censorship, combined with the discarding of established pandemic handling plans, suggests that the primary objective has been to frighten the population into compliance.**

**The Target Populations and the Precautionary Principle**

**The aggressive push to administer these injections to the entire population, including those who are not at risk, violates fundamental public health principles. If these measures were truly about health, they would be targeted only at those who stand to benefit most. Instead, injections are being pushed on children—who are not vulnerable to the virus—pregnant women, and individuals who have already recovered and possess natural immunity. The "precautionary principle" should dictate that we do not administer these products to healthy young people when the safety profile is so volatile and the long-term effects are**

**unknown. The current situation suggests that the vaccines are not a public health measure, but rather a tool used in a broader campaign of psychological and economic warfare.**

**Conclusion and Call to Action**

**The presence of "hot lots" and the extreme variability in vaccine toxicity constitute a violation of drug acts regarding adulterated products. There is an urgent need for an immediate halt to the COVID-19 vaccination program until these discrepancies are transparently investigated and clarified for the public. Because political institutions and the media are unlikely to act without significant pressure, citizens are encouraged to take a stand. This includes contacting representatives, demanding action from public prosecutors, and sharing information with those who have been misled by the media. Ultimately, reclaiming personal autonomy and returning to "normal" requires a collective refusal to comply with policies that are not based on transparent or honest health data. Taking courage and saying "I do not comply" is the essential first step in addressing what is characterized as a global crime.**

**To better understand the manufacturing of these vaccines, imagine a high-precision clockmaker who produces thousands of identical watches. If most of the watches keep perfect time, but every tenth watch is designed to explode, it is impossible to claim that the explosions are just a minor "mishap" in the assembly line of such a skilled professional.**

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